Amendments to the claims:

This listing of claims replaces all prior versions and listings of claims in the application.

Listing of Claims:

- 1. (Currently amended) An isolated peptide fragment of a natural cytotoxicity receptor of an NK cell, comprising a linker peptide connecting the extracellular domain of the receptor to the transmembrane portion of the receptor, wherein the isolated peptide fragment is about 10-100 amino acid residues in length and wherein said peptide fragment exhibiting exhibits at least one activity selected from binding to a viral infected cell or binding to a tumor cell.
- (Previously presented) The peptide fragment of claim 1 comprising at least one glycosylated residue.
- (Previously presented) The peptide fragment of claim 1 wherein the natural cytotoxicity receptor of the NK cell is selected from NKp46 and NKp44.
- 4. (Currently amended) The isolated peptide fragment of the human NKp46 receptor according to claim 3 comprising the amino acid sequence as set forth in SEQ ID NO:3, or an analog thereof, the peptide having the ability to bind to target cells selected from viral-infected cells and tumor cells, an active fragment, an isoform, an analog or a derivative thereof, with the proviso that said peptide is other than SEQ ID NOS:1 and 2.
- (Previously presented) The peptide fragment of claim 4 wherein the target cell is of a warm-blooded vertebrate.
- (Previously presented) The peptide fragment of claim 5 wherein the target cell is of human origin.
- (Previously presented) The peptide fragment of claim 4 comprising a minimal epitope of NKp46 receptor having ability to bind to viral-infected cells

- (Previously presented) The peptide fragment of claim 7 comprising a glycosylated residue corresponding to threonine at position 225 of isoform a of the human NKp46 receptor.
- (Previously presented) The peptide of claim 7 wherein the glycosylated residue comprises sialic acid.
- (Currently amended) The peptide fragment of claim 4 comprising from about 40-25 to 100-75 amino acids.
- 11. (Previously presented) The peptide fragment of claim 4 comprising from about 30 to 60 amino acids
- 12. 19. (Withdrawn)
- 20. (Currently amended) A fusion protein comprising a peptide wherein said peptide is-an isolated peptide fragment of a natural cytotoxicity receptor of an NK cell, and further comprising a molecule selected from an immunoglobulin (Ig) molecule or a fragment thereof, and a cytotoxic substance; the peptide fragment comprising the—a_linker peptide connecting the extracellular domain of the receptor to the transmembrane portion of the receptor, wherein the peptide fragment is about 10-100 amino acid residues in length; wherein said fusion protein comprising said peptide fragment exhibiting exhibits at least one activity selected from binding to a viral infected cell or binding to a tumor cell; and wherein said fusion protein is other than the fusion proteins of SEQ ID NOS:13-1816.
- (Original) The fusion protein of claim 20 manufactured by recombinant DNA technology or chemical synthesis.
- 22. (Currently amended) The fusion protein of claim 21-20 comprising a-the peptide fragment covalently conjugated to a molecule selected from an immunoglobulin (Ig) molecule or a fragment thereof, and a cytotoxic substance.

- 23. (Currently amended) The fusion protein of claim 22 wherein the peptide fragment is covalently conjugated to the Fc fragment of said immunoglobulin molecule.
- 24. (Currently amended) A pharmaceutical composition comprising as an active ingredient a peptide wherein said peptide is an isolated fragment of a natural cytotoxicity receptor of an NK cell, comprising the a_linker peptide connecting the extracellular domain of the receptor to the transmembrane portion of the receptor, wherein the isolated peptide fragment is about 10-100 amino acid residues in length and wherein said peptide fragment exhibiting-exhibits at least one activity selected from binding to a viral infected cell or binding to a tumor cell.
- 25. (Original) The pharmaceutical composition of claim 24 further comprising pharmaceutically acceptable diluents, carriers or excipients.
- (Previously presented) A pharmaceutical composition comprising as an active ingredient a fusion protein according to claim 20.
- 27. (Original) The pharmaceutical composition of claim 26 further comprising pharmaceutically acceptable diluents, carriers or excipients.
- 28. 33. (Canceled)
- 34. 46. (Withdrawn)
- 47. (Currently amended) A variant polypeptide comprising NKp46 receptor polypeptide or an active fragment thereof having at least a single amino acid substitution in an epitope required for the recognition of viral-infected cells or tumor cells, wherein the epitope is in the proximal domain of the NKp46 receptor.
- 48. (Currently amended) The variant polypeptide of claim 47, wherein the single amino acid substitution of NKp46 isoform a is Threonine 225 replaced by an amino acid residue selected from the group consisting of Serine, Alanine and Asparagine.